## Amendments to the claims

This listing of claims will replace all prior versions, and listings, of claims in the application:

- 1. (Original): A method of treating or preventing IBD in a mammal; comprising, administering a therapeutically effective amount of LXR agonist, or a pharmaceutically acceptable salt, solvate, or physiologically functional derivative thereof.
- 2. (Original): The method of claim 1 in which IBD is selected from the group consisting of Crohn's disease, ulcerative colitis, and inflammatory colitis caused by bacteria, ischemia, radiation, drugs or chemical substances.
- 3. (Currently amended): The method according to claim 1 or 2, wherein the LXR agonist is a compound of formula (II):

$$X \longrightarrow (CR^1R^2)_p \longrightarrow Z \longrightarrow (CH_2)_n \longrightarrow N \longrightarrow (CHR^4)_q \longrightarrow (CH$$

wherein:

X is OH or NH<sub>2</sub>;

p is 0-6;

each  $R^1$  and  $R^2$  are the same or different and are each independently selected from the group consisting of H,  $C_{1-8}$ alkyl,  $C_{1-8}$ alkoxy and  $C_{1-8}$ thioalkyl;

Z is CH or N;

when Z is CH, k is 0-4;

when Z is N, k is 0-3;

each  $R^3$  is the same or different and is independently selected from the group consisting of halo, -OH,  $C_{1-8}$ alkyl,  $C_{2-8}$ alkenyl,  $C_{1-8}$ alkoxy,  $C_{2-8}$ alkenyloxy,

 $-S(O)_aR^6, -NR^7R^8, -COR^6, COOR^6, R^{10}COOR^6, OR^{10}COOR^6, CONR^7R^8, -OC(O)R^9, \\$ 

-R<sup>10</sup>NR<sup>7</sup>R<sup>8</sup>, -OR<sup>10</sup>NR<sup>7</sup>R<sup>8</sup>, 5-6 membered heterocycle, nitro, and cyano;

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a is 0, 1 or 2;

 $R^6$  is selected from the group consisting of H,  $C_{1-8}$ alkyl,  $C_{1-8}$ alkoxy and  $C_{2-8}$ alkenyl;

each  $R^7$  and  $R^8$  are the same or different and are each independently selected from the group consisting of H,  $C_{1-8}$ alkyl,  $C_{2-8}$ alkenyl,

C<sub>3-8</sub>alkynyl;

 $R^9$  is selected from the group consisting of H,  $C_{1-8}$ alkyl and -NR<sup>7</sup>R<sup>8</sup>;  $R^{10}$  is  $C_{1-8}$ alkyl;

n is 2-8;

q is 0 or 1;

R<sup>4</sup> is selected from the group consisting of H, C<sub>1-8</sub>alkyl, C<sub>1-8</sub>alkenyl, and alkenyloxy;

Ring A is selected from the group consisting of C<sub>3-8</sub>cycloalkyl, aryl, 4-8 membered heterocycle, and 5-6 membered heteroaryl;

each ring B is the same or different and is independently selected from the group consisting of C<sub>3-8</sub>cycloalkyl and aryl.

4. (Original): The method according to claim 3, in which the LXR agonist is the compound of formula (IIa)

(IIa)

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5. (Currently amended): The method according to claim 1 or 2, wherein the LXR agonist is a compound of compounds of formula (I):

$$X^{1} \xrightarrow{X^{2}} X^{3}$$

$$R^{1} \xrightarrow{Ar-Y} X^{6}$$

$$X^{4} \xrightarrow{X^{5}} X^{6}$$

$$(I)$$

wherein:

Ar represents an aryl group;  $R^1$  is -OH, -O-( $C_1$ - $C_7$ )alkyl, -OC(O)-( $C_1$ - $C_7$ )alkyl, -O-( $C_1$ - $C_7$ )heteroalkyl, -OC(O)- ( $C_1$ - $C_7$ )heteroalkyl, -CO<sub>2</sub>H, -NH<sub>2</sub>, -NH( $C_1$ - $C_7$ )alkyl, -N(( $C_1$ - $C_7$ )alkyl)<sub>2</sub> or -NH-S(O)<sub>2</sub>-( $C_1$ - $C_5$ )alkyl;

 $R^2$  is  $(C_1-C_7)$ alkyl,  $(C_1-C_7)$ heteroalkyl, aryl and aryl $(C_1-C_7)$ alkyl;

- $X^1$ ,  $X^2$ ,  $X^3$ ,  $X^4$ ,  $X^5$  and  $X^6$  are each independently H, (C<sub>1</sub>-C<sub>5</sub>)alkyl, (C<sub>1</sub>-C<sub>5</sub>)hetroalkyl, F or Cl, with the proviso that no more than three of  $X^1$  through  $X^6$  are H, (C<sub>1</sub>-C<sub>5</sub>)alkyl or (C<sub>1</sub>-C<sub>5</sub>)heteroalkyl; and
- Y is  $-N(R^{12})S(O)_{m^-}$ ,  $-N(R^{12})S(O)_{m}N(R^{13})$ -,  $-N(R^{12})C(O)$ -,  $-N(R^{12})C(O)N(R^{13})$ -,  $-N(R^{12})C(S)$  or  $-N(R^{12})C(O)O$ -, wherein R12 and R13 are each independently hydrogen,  $(C_1-C_7)$ aryl,  $(C_1-C_7)$ heteroalkyl, aryl and  $aryl(C_1-C_7)$ alkyl, and optionally when Y is  $-N(R^{12})S(O)_{m^-}$  or  $-N(R^{12})S(O)_{m}N(R^{13})$ -,  $R^{12}$  forms a five, six or seven-membered ring fused to Ar or to  $R^2$  through covalent attachment to Ar or  $R^2$ , respectively. In the above Y groups, the subscript m is an integer of from 1 to 2.
- 6. (Original): The method according to claim 5, in which the LXR agonist is the compound of formula Ia

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